

Clinixel Life Sciences Pvt Ltd - Trusted CRO in India

About Us

Clinixel is a full-service Contract Research Organization with headquarters in **Mumbai, India**, and a presence in **New Jersey, USA**. We proudly serve emerging biotechnology, mid-size pharmaceutical, and medical device companies across **India, the USA, Europe, China, Africa, Australia, and Israel**, thanks to our membership in the AICROS global CRO alliance.

Led by industry experts—physicians, pharmacists, and doctorate-level professionals—Clinixel brings decades of collective experience in clinical development, regulatory affairs, pharmacovigilance, and medical writing



What We Offer — Comprehensive CRO Services

Clinical Trials & Project Management

We deliver end-to-end trial services, from feasibility and regulatory submissions to site selection, logistics, recruitment, monitoring, and CSR generation—covering Phases I–IV, biosimilars, new chemical entities, and medical devices.

Global Clinical Development

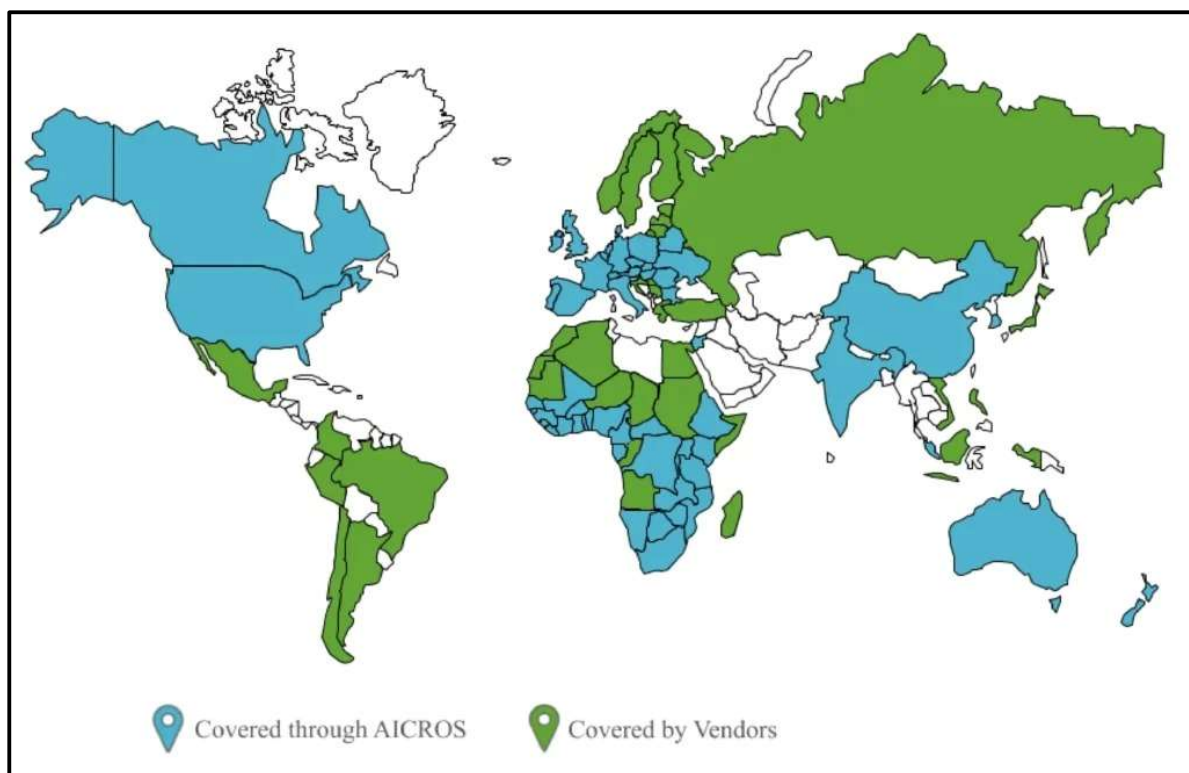
Through our AICROS network, we facilitate multinational trials with harmonized submissions, logistics handling (IP and sample import/export), centralized data management, and unified client communication—delivering seamless project execution across **80+ countries**.

Pharmacovigilance

Our services span ICSR processing, aggregate reporting (DSUR, PBRER), literature monitoring, signal management, and risk management planning—ensuring proactive safety oversight Biomed Consult.

Medical Writing, Regulatory Affairs, Biostatistics & Data Management

Clinexel's skilled team develops high-quality regulatory dossiers, clinical study reports, statistical analyses, and secure data handling systems tailored to your study needs.



Note: We have demonstrated success in conducting clinical trials across all 4 phases (phase I-IV) including First in Man, in Healthy Volunteers and in Patient's.

Our Strengths

- **In-House Expertise:** We rely on dedicated full-time staff—CRAs, medical monitors, project managers, writers—ensuring consistency, quality, and alignment with global standards.
- **Regulatory & Quality Compliance:** Deep experience with ICH-GCP, EPA-style regulations, CDSCO compliance, and adherence to global trial conduct standards ensures seamless regulatory alignment.
- **Proven Efficiency in Trial Setup & Logistics:** We excel in realistic feasibility projections, recruitment timelines, cost estimation, site readiness, and supply chain management—optimized for timely execution.
- **Accolades & Recognition:** Clinexel has been honored as the **Best Full-Service CRO — South Asia 2023** and **Best Medical Writing CRO — Asia 2024**, affirming our leadership in quality and innovation.

Why Clinexel?

- A **trusted global partner** with a local touch.
- **Experienced leadership** rooted in clinical and safety excellence.
- **Scalable services** for trials of any phase or therapeutic area.
- **Efficient operations**, from regulatory planning to final reporting.
- **Proven track record**, recognized by industry awards.

Contact Us

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